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Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims:

1. (Currently Amended) A method to manufacture composite polymer electrolyte membranes coated with inorganic thin films for fuel cells, ~~characterized to obtain wherein~~ composite membranes are obtained by coating the surface of polymer electrolyte membranes with inorganic thin films ~~using a plasma-enhanced chemical vapor deposition method or a reactive sputtering method.~~
2. (Currently Amended) The method ~~according to~~ of claim 1 ~~[[,]]~~ wherein the inorganic materials of said inorganic thin film are chosen one or more from the group comprising silicon oxide (SiO_2), titanium oxide (TiO_2), zirconium oxide (ZrO_2), zirconium phosphate ($\text{Zr}(\text{HPO}_4)_2$), zeolite, silicalite, and aluminum oxide (Al_2O_3).
3. (Currently Amended) The method ~~according to~~ of claim 1 ~~[[,]]~~ wherein said polymer electrolyte membranes are ~~[[,]]~~ perfluorosulfonic acid membranes such as Nafion[®] membrane, Dow membrane, Flemion membrane, Aciplex membrane, BAM, or Gore-select membrane; electrolyte membranes made of proton conducting hydrocarbon materials such as sulfonic polysulfonide, sulfonic polyethylene, sulfonic polypropylene, sulfonic polystyrene, sulfonic polyphenol formaldehyde, polystyrene divinylbenzene sulfonic acid, sulfonic polybenzimidazole, sulfonic polyamide, or sulfonic polyether-ether ketone; or electrolyte membranes made of proton conducting fluorine materials such as sulfonic polyvinylidene fluoride, sulfonic polytetrafluoroethylene, or fluoroc ethylene propylene.

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4. (Currently Amended) The method ~~according to claim 1, of claim 19~~ wherein said PECVD method uses reactants ~~being comprising~~ one or more monomers chosen from the group of organic metal compounds containing aluminum, titanium, silicon, and zirconium in conjunction with one or more gases out of the group of oxygen, nitrogen, hydrogen, steam, and argon.

5. (Currently Amended) The method ~~according to claim 1 of claim 4~~ wherein said organic metal compounds are one or more chosen from the group comprising trimethyl disiloxanes (TMDSO), hexamethyl disiloxane (HMDSO), hexamethyl disilane, tetraethyl orthosilicate (TEOS), tetramethyl orthosilicate, tetrabutyl orthosilicate, tetra-isopropyl orthosilicate, aluminium methoxide, aluminium ethoxide, aluminium butoxide, aluminium isopropoxide, titanium ethoxide, titanium methoxide, titanium butoxide, titanium isopropoxide, zirconium ethoxide, and zirconium butoxide.

6. (Currently Amended) The method ~~according to of claim 2,~~ wherein said organic metal compounds are one or more chosen from the group comprising trimethyl disiloxanes (TMDSO), hexamethyl disiloxane (HMDSO), hexamethyl disilane, tetraethyl orthosilicate (TEOS), tetramethyl orthosilicate, tetrabutyl orthosilicate, tetra-isopropyl orthosilicate, aluminium methoxide, aluminium ethoxide, aluminium butoxide, aluminium isopropoxide, titanium ethoxide, titanium methoxide, titanium butoxide, titanium isopropoxide, zirconium ethoxide, and zirconium butoxide.

7. (Currently Amended) The method ~~according to claim 1, of claim 20~~ wherein said reactive sputtering process is characterized to use a 99 % or higher pure metal target such as Si, SiO₂, SiNH, Al, Zr, or Ti, and to maintain its initial pressure at a high vacuum range of $1.0 \cdot 10^{-3}$

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torr to 1.0×10^{-6} torr.

8. (Currently Amended) The method ~~according to claim 1, of claim 19~~ wherein said PECVD method ~~or reactive sputtering method~~ is characterized to have a microwave power at the range of 10 watts to 500 watts.

9. (Currently Amended) The method ~~according to claim 1, of claim 19~~ wherein the reaction chamber pressure of said PECVD method ~~or reactive sputtering method~~ is in the range of 1.0 to 1000 millitorr.

10. (Currently Amended) The method ~~according to claim 1, of claim 19~~ wherein the argon pre-treatment electromagnetic wave power of said PECVD method ~~or reactive sputtering method~~ is in the range of 10 watts to 500 watts.

11. (Currently Amended) The method ~~according to claim 1, of claim 19~~ wherein the argon pre-treatment pressure of said PECVD method is in the range of 1.0 to 500 millitorr.

12. (Currently Amended) The method ~~according to claim 1, of claim 19~~ wherein the reaction gas pressure in the chamber of said PECVD process is in the range of 10 to 500 millitorr.

13. (Currently Amended) The method ~~according to claim 1, of claim 1~~ wherein the thickness of said inorganic films is in the range of 1.0 to 500 nm.

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14. (Currently Amended) The method ~~according to claim 1, of claim 1~~ wherein said manufacturing method further comprises a step of coating the surface of electrolyte membrane with a proton-conducting ionomer solution, after coating said inorganic film on the said membrane surface, so as to enhance contact with the electrodes during manufacturing membrane-electrode assembly.
15. (Withdrawn) A composite polymer electrolyte membrane coated with inorganic thin films for fuel cells manufactured according to claim 1.
16. (Withdrawn) An MEA employing the composite polymer electrolyte membranes coated with inorganic thin films manufactured according to claim 1.
17. (Original) A method of manufacturing an MEA including a process of coating catalysts for electrodes directly on the composite polymer electrolyte membranes coated with inorganic thin films manufactured according to claim 1.
18. (Withdrawn) A fuel cell employing the composite polymer electrolyte membranes coated with inorganic thin films or the MEA containing the said composite membrane manufactured according to claim 1.
19. (New) The method of claim 1 wherein the inorganic thin film is coated on the surface of the polymer electrolyte membrane using a plasma enhanced chemical vapor deposition (PECVD) method.
20. (New) The method of claim 1 wherein the inorganic thin film is coated on the surface of the polymer electrolyte membrane using a reactive sputtering method.

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21. (New) The method of claim 20 wherein said reactive sputtering method is characterized to have a microwave power at the range of 10 watts to 500 watts.
22. (New) The method of claim 20 wherein the reaction chamber pressure of said reactive sputtering method is in the range of 1.0 to 1000 millitorr.
23. (New) The method of claim 20 wherein the argon pre-treatment electromagnetic wave power of said reactive sputtering method is in the range of 10 watts to 500 watts.